Version 1.2, Nov 2002

OLIN-WILMINGTON LEVEL I DATA QUALITY EVALUATION STANDARD OPERATING PROCEDURE AND CHECKLIST ICP METALS BY METHOD 6010B/200.7

Reviewer/Date	Tige CUNNINGhan	- 6/18/10
Sr. Review/Date	Tige CUNDINGHAM	4/25/10
Lab Report # 30	0-26874-1	1 1
Project # 60	7100016-12	

OC-GW-IGR ONLY for Dissolud Fe **Laboratory Deliverable Requirements** 1.0 1.1 Laboratory Information: Was all of the following provided in the laboratory report? Yes Vo No N/A Check items received. Address

Project ID

Phone # Sample identification – Field and Laboratory

Address

Client Contact

(IDs must be cross-referenced) ☑ Name of Laboratory Client Information: ACTION: If no, contact lab for submission of missing or illegible information. Yes [No [] N/A [] 1.2 Laboratory Report Certification Statement Does the laboratory report include a completed Analytical Report Certification in the required format? ACTION: If no, contact lab for submission of missing certification or certification with correct format. Yes No No N/A Comments: 1.3 Laboratory Case Narrative: Narrative serves as an exception report for the project and method QA/QC performance. Narrative includes an explanation of each discrepancy on the Certification Statement. ACTION: If no, contact lab for submission of missing or illegible information. 1.4 Chain of Custody (COC) copy present with all documentation completed NOTE: Olin receives and maintains the original COC.

ACTION: If no, contact lab for submission of copy of completed COC.

	1.5 Sample	Receipt	Information (Cooler Rece	eipt Form	present?):		Yes [V]	No []	N/A [_]	Comments:		
, i	Were each o		lowing tasks of	ompleted and	recorded u	ipon receipt of the	sample(s)						
Sam	ole temperature	confirm	ed: must be 1°	– 10° C. (If sa	amples were	e sent by courier and	d delivered o	on the same o	lay as colle	ction, tempera	ture requirement of	loes not apply).	
Cont	ainer type note	d ⊡ san	nple condition	observed 🗹 j	pH verified	(where applicable)	Field and	d lab IDs cro	ss reference	ed			
ACTIC	N: If no, conta	act lab fo	or submission o	of missing or in	ncomplete d	locumentation.							
	1.5.1	Were a	all samples del	ivered to the la	aboratory w	ithout breakage?		Yes	No [_]	N/A [_]	Comments:		
	1.5.2	with sa	the Cooler Reample receipt, astances affect	condition of th	e samples,	ntive indicate other analytical problems	problems or special	Yes []	No 🗹	N/A [_]	Comments:		
			Section: Wa for each samp		following	requirements suppl	lied in the	Yes	No []	N/A [_]	Comments:		
Fie Clea	ld ID and Lab an-up method rix	ID M/A	Date and Analysis Target analy	d time collecte method ytes and conce	ed ntrations	Analyst Initials Preparation m	nethod Uni	Dilution Date of p ts (soils must	Factor reparation/o	extraction/dige	noisture or solids estion clean-up an	MA PRODUCE RE	eporting lim ere applicab
ACTIO	ON: If no, con	tact lab f	or submission	of missing or i	incomplete	information.							
			nation: Was each sample ba		following	information supplie	ed in the	Yes [V]	No []	N/A []	Comments:		

/		1				and the second second	
Metl	nod blank results	s LCS recoverie	MS/MSD recoveries and RPDs Laboratory	duplicate res	ults (where	applicable)	
ACTIO	N: If no, contac	ct lab for submission	of missing or incomplete information.				
	and the second s		erencis misera dicessi — Produce i sissele editare i → di prese editioni viva di Producti (1974				
2.0	Holding Tim	es					
					/		
Have an	ny technical ho exceeded? Ho water and soil.	olding time for metals	ned from date of collection to date of analysis, been is 180 days from sample collection to analysis for both	Yes [_]	No 🗹	N/A]	Comments:
NOTE:	List samples tha	at exceed hold time w	rith # of days exceeded on checklist	18.			
nord.	Dist sumples the	at exceed fiold time (in a crange exceeded on encountry				
ACTIO	N: If technical	holding times are e	acceeded, qualify all positive results (J) and non-detects				
110110		_	ng time) reject (R) all non-detect results.				
	A. S.						
3.0	Laboratory	Method					
	Article Harrison Control	- 1 - 42/10 - 12/10		2			
	3.1 Was	the correct laborate	ory method used?	Yes [V]	No []	N/A [_]	Comments:
	Wate	er Digestion	3005A or 3010A or 3020A				
		Digestion	3050B				
	Meta		6010B or 200.7				
comp		quested method. C	y to provide justification for method change ontact senior chemist to inform Client of change				
	3.2 Are	the practical quant SOW □ QAP	itation limits the same as those specified by the P	Yes 🔼	No []	N/A []	Comments:
NOT	E: Verify that th	e reported metals ma	tch the target list specified on the COC.				
	28			AD:			
6010.de	oc					*	

		If no, evaluate variation with respect to sample matrix, preparation, dilution, c. If sample PQL is indeterminate, contact lab for explanation.				
	3.3	Are results present for each sample in the SDG?	Yes L	No [_]	N/A []	Comments:
ACTIO	ON: If no	o, check Request for Analysis to verify if method was ordered and COC to verify that it				
	3.4	If dilutions were required, were dilution factors reported?	Yes 🔄	No [_]	N/A []	Comments:
ACTIO	ON: If no	o, contact the lab for submission.				
4.0	Meth	od Blanks				
	4.1	Is the Method Blank Summary present?	Yes [V]	No []	N/A []	Comments:
ACT	ΓΙΟΝ:	If no, call the laboratory for submission of missing data.				
	4.2	Frequency of Analysis: Was a method blank analyzed for each digestion batch of < 20 field samples?	Yes 🗹	No [_]	N/A [_]	Comments:
		If no, contact laboratory for justification. Consult senior chemist for action rate non-compliance.		7		
	4.3	Is the method blank less than the PQLs for all target elements?	Van I	Nat 1	NI/A F 1	Comments:
NOT sam		DEP requires the method blank to be matrix matched and digested with the	res [<u>v</u>]	NO []	N/A [_]	Comments.
	4.4 the fol	Do any method blanks have positive results for metals? Qualify data according to lowing:	Yes [_]	No []	N/A []	Comments:

If the sample concentration is $\leq 5 \times$ blank value, flag sample result non-detect "U" at the	
PQL or the concentration reported if greater than the PQL.	

If the sample concentration is $> 5 \times$ blank value, no qualification is needed.

ACTION: For any blank with positive results, list all contaminants for each method blank including the concentration detected and the flagging level (flagging level = 5x the blank value) and the associated samples and qualifiers.

= 5x th	e blank	value) and the associated samples and qualifiers.				
5.0	Labo	ratory Control Standard				
	5.1	Was a laboratory control standard run with each analytical batch of 20 samples or less?	Yes 🔽	No []	N/A []	Comments:
ACT	ION:	Ill target, second source LCS is required by MADEP. Call laboratory for LCS form submittal. If data are not available, use judgement to evaluate data accuracy associated with that batch.		/		
	5.2	Is a LCS Summary Form present?	Yes [No []	N/A []	Comments:
ACT	ION:	If no, contact lab for resubmission of missing data.			/	
	5.3 Samp	Is the recovery of any analyte outside of MADEP control limits? MADEP le Type 80 120	Yes []	No [V]	N/A [_]	Comments:
	water	80-120				
withi	in the l detects	within Lab generated limits If recovery is above the upper limit, qualify all positive sample results batch as (J). If recovery is below the lower limit, qualify all positive and results within the batch as (J). If LCS recovery is <30%, positive and nonsare rejected (R).				

Comments:

Yes | | No | | N/A |

6.0 **Matrix Spikes**

Matrix spikes may be collected at different frequencies based on monthly, quarterly, or task specific schedules. Confirm spike requirements for each set with the senior chemist.

Were project-specific MS/MSDs collected? List project samples that were 6.1

ACTION: If no, contact senior chemist to see if any were specified.

6.2 Is the Matrix Spike/Matrix Spike Duplicate Recovery Form present?

NOTE: A full target, second source MS/MSD is required by MADEP.

ACTION: If any matrix spike data are missing, call lab for resubmission.

Were matrix spikes analyzed as indicated on the COC and project 6.3 schedule?

ACTION: If any matrix spike data are missing, call lab for resubmission. If none, no qualification is needed. Narrate non-compliance.

Are any metal spike recoveries outside of the QC limits? 6.4

	MADEP	QAPP	
Sample Type	% Rec	% Rec	Method
Water	75-125	N/A	6010B
Water	N/A	70-130	200.7
Soil	75-125	75-125	6010B

NOTE: %R = (SSR-SR) x 100% SA

Where: SSR = Spiked sample result SR = Sample result

SA = Spike added

NOTE: If dilutions are required due to high sample concentrations (> 4X spike), the data are evaluated, but no flags are applied.

NOTE: If only one of the recoveries for an MS/MSD pair is outside of the control limits, no qualification is necessary. Use professional judgment for the MS/MSD flags.

ACTION: MS/MSD flags only apply to the sample spiked. If the recoveries of the MS and MSD exceed the upper control limit, qualify positive results as estimated (J). If the recoveries of the MS and MSD are lower than the lower control limit, qualify positive results and non-detects (J).

6.5 Are any RPDs for MS/MSD recoveries outside of the QC limits?

Yes No N/A Comments

NOTE: RPD = S-D x 100% (S+D)/2

Where: S = MS sample result D = MSD sample result

NOTE: If dilutions are required due to high sample concentrations, the data are evaluated, but no flags are applied.

ACTION: If the RPD exceeds the control limit, qualify positive results and non-detects (J).

7.0 Laboratory Duplicate

7.1 Was a laboratory duplicate sample analyzed? If so, is the Laboratory Yes No NA Duplicate Sample Form present?

NOTE: MADEP refers to this sample as a "matrix duplicate".

ACTION: If not analyzed, qualification is not needed. If data is missing, contact laboratory for resubmission of report. Narrate non-compliance.

7.2 Is the RPD between the result for the laboratory duplicate sample and the result for the parent sample outside of the QA/QC limits?

Yes No NA Comments

MADEP Laboratory Duplicate Sample RPD Criteria:	QAPP RPD
For aqueous results > $5 \times$ RL, RPD must be $\pm 20\%$	20
For aqueous results < 5× RL, RPD must be < RL	20
For soil/sediment results > $5 \times$ RL, RPD must be $\pm 35\%$	20
For soil/sediment results < $5 \times RL$, RPD must be $\leq 2 \times RL$	20

ACTION: If the RPD exceeds the limits, qualify both positive results and non-detects as estimated and flag them J. Narrate non-compliance

8.0 Sampling Accuracy

The majority of ground water samples are collected directly from a tap, process stream, or with dedicated tubing. Rinse blanks will not be collected.

- **8.1** Were rinsate blanks collected? Prior to evaluating rinsate blanks, obtain a list of the associated samples from the senior chemist.
- **8.2** Do any rinsate blanks have positive results?

NOTE: MADEP does not require the collection of rinsate blanks.

ACTION: Evaluate rinsate results against blank results to determine if contaminant may be laboratory-derived. If results are not lab-related, qualify according to below.

If the sample concentration is $< 5 \times$ blank value, flag sample result non-detect "U" at the PQL or the concentration reported if greater than the PQL.

If the sample concentration is $> 5 \times$ blank value, no qualification is needed.

9.0 Field Duplicates

9.1 Were field duplicate samples collected? Obtain a list of samples and their associated field duplicates.

Yes No N/A Comments

9.2 Were field duplicates collected per the required frequency?	Yes []	No No	N/A [_]	Comments:			
SOW □ QAPP (1 per 10) □ MADEP Option 1 (1 per 20) □ MADEP Option 3 (1 per 10) □							
9.3 Was the RPD \leq 50% for soils or waters? Calculate the RPD for all results and attach to this review.	Yes []	No [_]	N/A	Comments:			
ACTION : RPD must be ≤50% for soil and water. Qualify data (J) for both sample results if	ACTION: RPD must be ≤50% for soil and water. Qualify data (J) for both sample results if the RPD exceeds 50%.						
10.0 Special QA/QC							
10.1 Were both total and dissolved metals analysis performed? If so, the dissolved metal concentration should not exceed that of the total metal.	Yes []	No I	N/A [_]	Comments:			
ACTION: If results for both total and dissolved are $\geq 5x$ the PQL and the dissolved concentration is 10% higher than the total, flag both results as estimated (J). If total and dissolved concentrations are less than $5x$ the PQL and the difference exceeds $2x$ the PQL, flag both results as estimated (J)							

10.0	Application	of Validation	Oualifiers
10.0	T X D PART CELLIO AL	OI THIIMMEION	O HIGHWAY CAR

Was any of the data qualified?

Yes [_]	No L	N/A []	Comments

If so, apply data qualifiers directly to the DQE copy of laboratory report and flag pages for entry in database.

REFERENCES

- LAW, 1999, "Final Quality Assurance Project Plan, Olin Wilmington Property, 51 Eames Street, Wilmington, MA", LAW Engineering and Environmental Services, Kennesaw, GA 30144. August 1999
- U.S. Environmental Protection Agency (USEPA), 1989. "Region 1 Laboratory Data Validation Functional Guidelines For Evaluating Inorganic Analyses"; Hazardous Site Evaluation Division; February 1989.
- MADEP, 2001. Massachusetts Department of Environmental Protection Bureau of Waste Site Cleanup, "Massachusetts Quality Assurance and Quality Control (QA/QC) Requirements." BWSC-CAM, Interim Final Draft, Revision No. 2, 5 October 2001.
- MADEP, 2001. Massachusetts Department of Environmental Protection Bureau of Waste Site Cleanup, "Quality Assurance and Quality Control Guidelines for Sampling, Data Evaluation and Reporting Activities," BWSC-CAM, Section VII, Public Comment Draft, Revision No. 0, 21 December 2001.